

**MALARIA**

# COURSE OBJECTIVES

- BASIC UNDERSTANDING OF MALARIA
  - EPIDEMIOLOGY
  - SYMPTOMS
  - DIAGNOSIS
  - TREATMENT
  - PREVENTION

# WHAT IS IT?

- A MOSQUITO-BORNE INFECTIOUS DISEASE OF THE TROPICS, SUBTROPICS, AND FRINGES OF TEMPERATE FORESTS .
  - FOUND IN LATIN AMERICA, CARIBBEAN, ASIA, AFRICA, EUROPE
- CAN BE EITHER ACUTE OR CHRONIC

# WHAT IS IT?

- CAUSED BY PROTOZOAN PARASITE  
GENUS *PLASMODIUM*
- FOUR SPECIES:
  - *P FALCIPARUM*
  - *P VIVAX*
  - *P OVALE*
  - *P MALARIAE*

# WHY THE CONCERN?

- MOST PREVALENT DISEASE IN THE WORLD
  - 2.1 BILLION LIVE IN MALARIOUS AREAS
  - 100-300 MILLION NEW CASES ANNUALLY
  - 1-3 MILLION DEATHS ANNUALLY
- POTENTIALLY LETHAL DISEASE
- SERIOUS THREAT TO MILITARY OPS



# HISTORY

- WWI NAVAL FORCES
  - 4,746 NEW HOSPITAL ADMISSIONS
  - 68,373 LOST MAN-DAYS
  - 7 DEATHS



# HISTORY

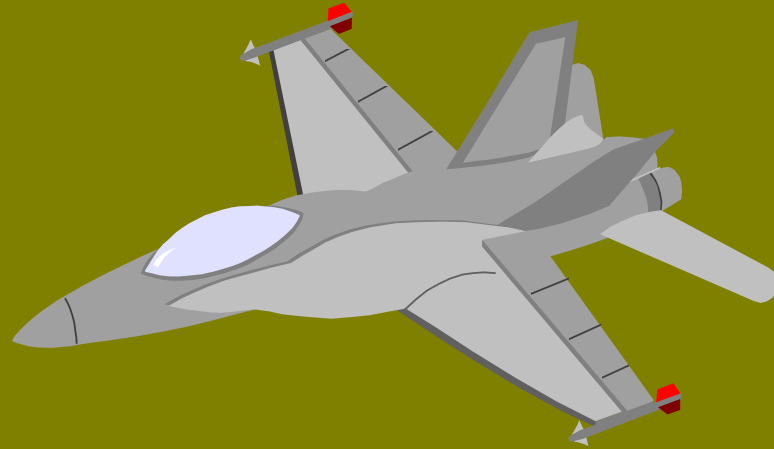
- WWII NAVAL FORCES
  - 111,675 NEW HOSPITAL ADMISSIONS
  - 3,310,800 LOST MAN-DAYS
  - 90 DEATHS
  - 5,332 (AVG) DAILY SICK LIST IN PACIFIC





# HISTORY

- VIETNAM WAR
  - 21,695 NEW ADMISSIONS
  - 187,478 LOST MAN-DAYS
  - 46 DEATHS



# TRANSMISSION

- MAN IS THE ONLY IMPORTANT RESERVOIR
- VECTOR IS FEMALE ANOPHELES MOSQUITO
  - **TEMPERATURE**
  - **RAINFALL**
  - ALTITUDE
  - TERRAIN

# TRANSMISSION

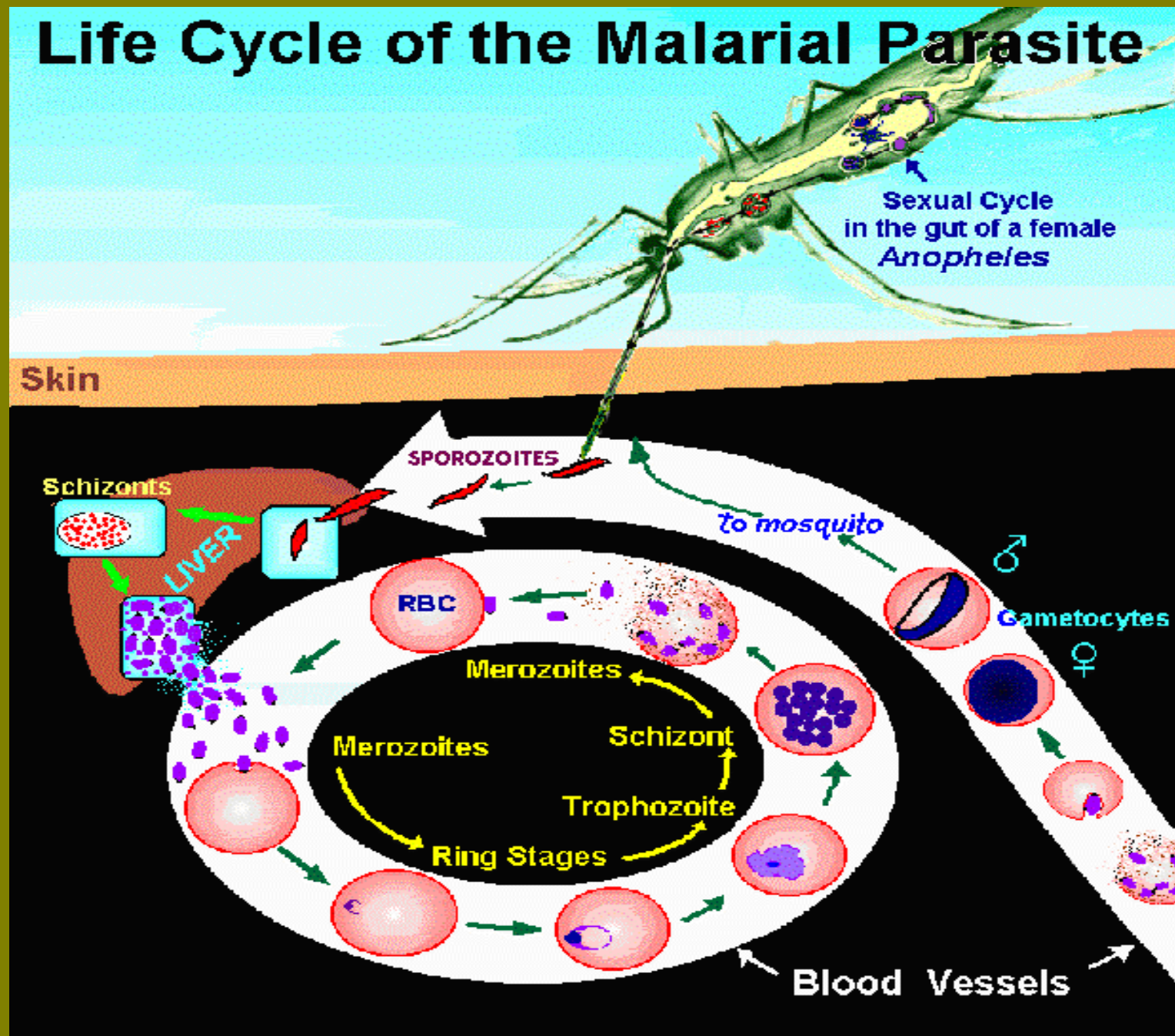
- MOSQUITO VECTOR:  
*ANOPHELES*
- BLOOD TRANSFUSION
- CONTAMINATED NEEDLE
- ORGAN TRANSF
- CONGENITAL



# SUSCEPTIBILITY

- UNIVERSAL SUSCEPTIBILITY
- NO ABSOLUTE IMMUNITY
  - PARTIAL IMMUNITY IN AREAS OF HIGH ENDEMICITY

# Life Cycle of the Malarial Parasite



# PATHOGENESIS

- RED BLOOD CELL DESTRUCTION
- IMMUNE COMPLEXES AND MEDIATORS
- CAPILLARY PERMEABILITY
- TISSUE HYPOXIA



# *PLASMODIUM SPECIES*

- *P. FALCIPARUM*
  - MOST SEVERE AND PREVALENT
  - 40-60% OF CASES
  - WIDESPREAD CHLOROQUINE RESISTANCE
  - INFECTS RBCs OF ALL AGES: HEAVY PARASITEMIA

# PLASMODIUM SPECIES

- P. VIVAX
  - 30-40% OF CASES
  - LIVER PHASE
  - INFECTS YOUNG RBCs: LESS SEVERE THAN FALCIPARUM
- OVALE
  - LIVER PHASE
  - INFECTS YOUNG RBCs
- MALARIAE
  - CAN PERSIST SUBCLINICALLY FOR EXTENDED PERIODS OF TIME
  - INFECTS OLD RBCs



# INCUBATION PERIOD

- |                 |          |
|-----------------|----------|
| • P. FALCIPARUM | 12 DAYS  |
| • P. VIVAX      | 14 DAYS* |
| • P. OVALE      | 14 DAYS* |
| • P. MALARIAE   | 30 DAYS  |

\* MAY BE 8 - 10 MONTHS OR  
LONGER FOR SOME STRAINS

# ACUTE SYMPTOMS

- CLASSICAL CYCLIC PAROXYSM:
  - COLD STAGE: CHILLS AND SHAKING
  - HOT STAGE: WARM, HEADACHE, VOMITING
  - SWEATING STAGE: WEAKNESS
  - FEEL WELL FOR PERIOD OF TIME, THEN CYCLE REPEATS ITSELF

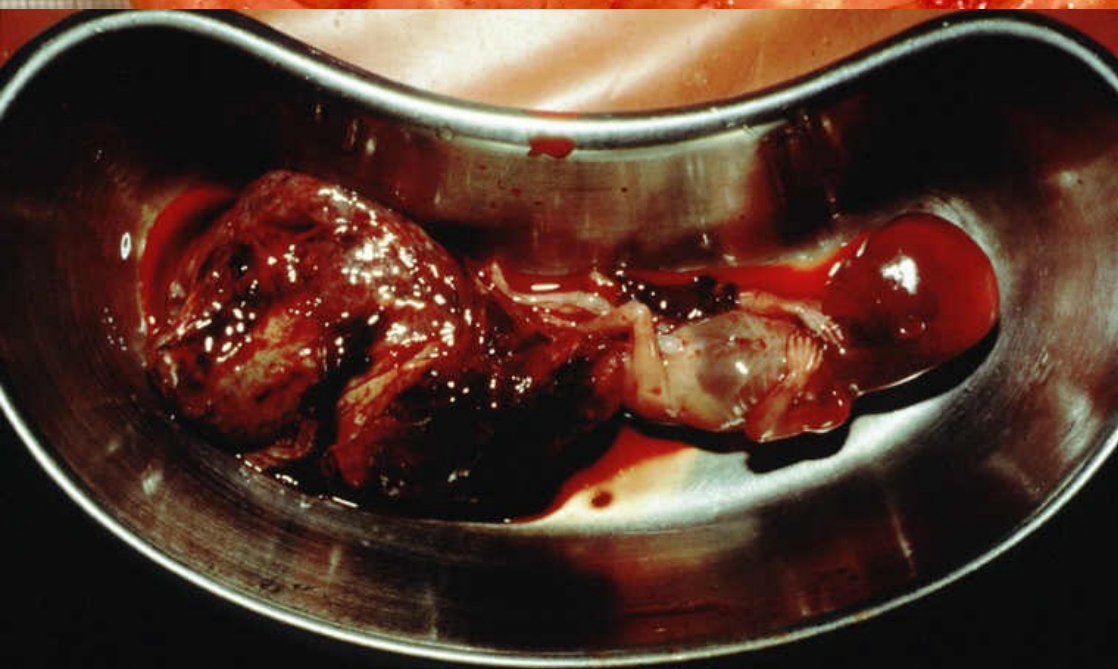
# PRESENTATION

- FEVER 97%
- CHILLS 97%
- HEADACHE 97%
- NAUSEA OR VOMITING 62%
- ABDOMINAL PAIN 56%
- MYALGIA 50%
- BACKACHE 9%
- DARK URINE 3%

# COMPLICATED MALARIA

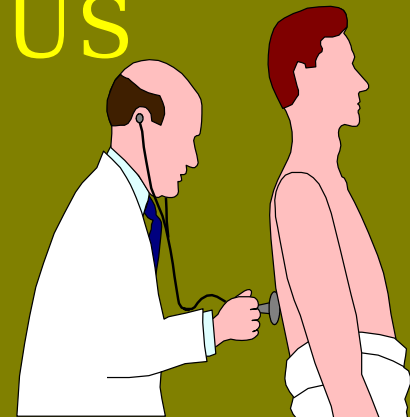
- HYPERPARISITEMIA: >3%
- HYPOGLYCEMIA: <60 MG/DL
- SEVERE ANEMIA:
  - HCT < 21% OR RAPIDLY FALLING HCT
- RENAL FAILURE
- HYPONATREMIA
- CEREBRAL MALARIA
- PROLONGED HYPOTHERMIA
- HIGH OUTPUT VOMITING OR DIARRHEA
- PREGNANCY





# SIGNS IN ACUTE INFECTION

- SLIGHTLY ILL TO IN DISTRESS
- ALERT TO UNCONSCIOUS
- FEVER



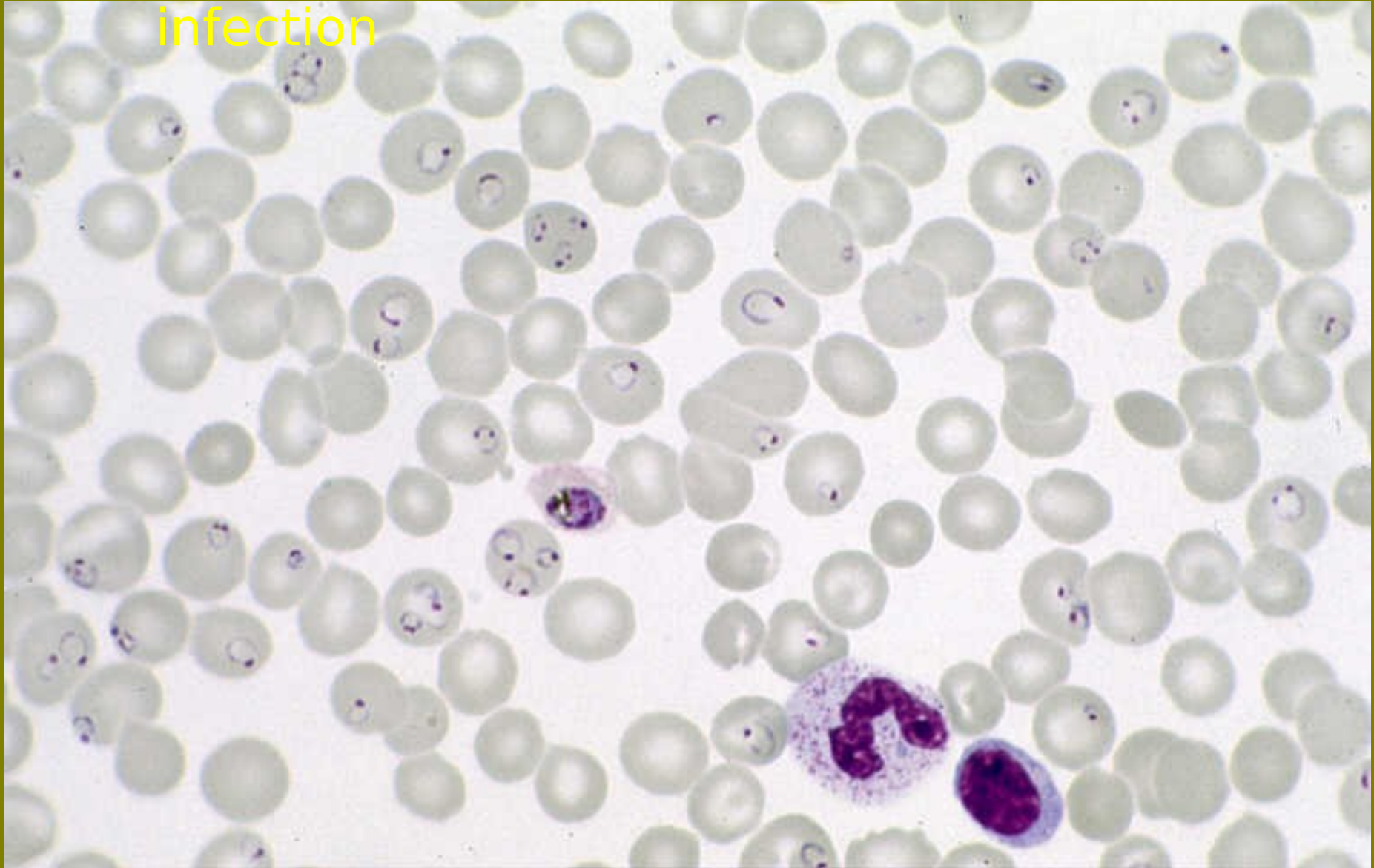


# DIAGNOSIS

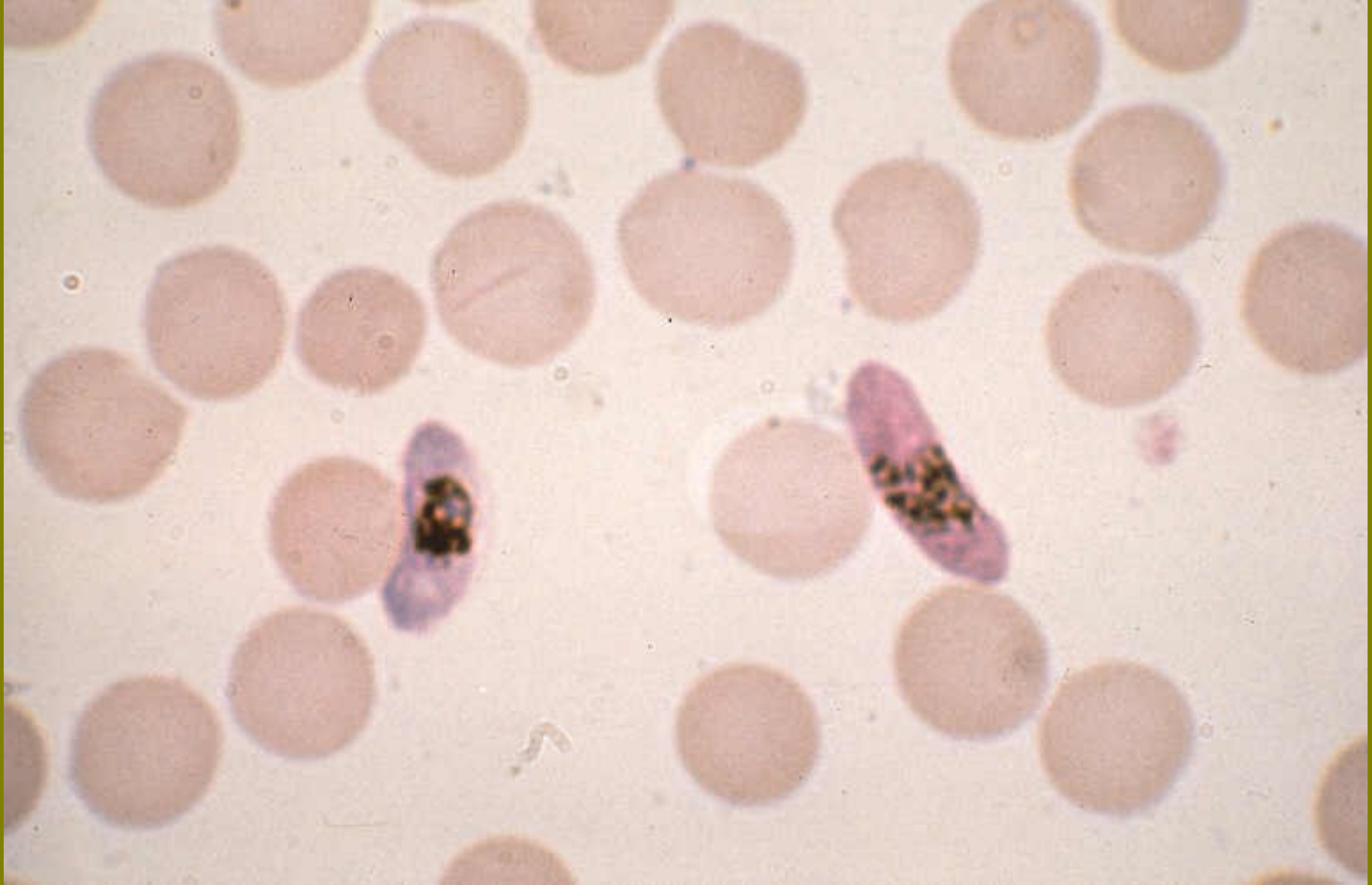
- ♦ **GOLD STANDARD: MULTIPLE THICK AND THIN SMEARS**
- ♦ **DIP STICK TESTS**
- ♦ **CBC**
  - **ANEMIA**
  - **LEUKOPENIA, OR LEUKOCYTOSIS**
  - **NO EOSINOPHILIA**



# Multiple RBC's with trophozoite stage of infection







Schizont stage of malarial infection

# TREATMENT

- CHLOROQUINE SENSITIVE INFECTIONS:

- CHLOROQUINE BASE 600 MG (2 TABS) P.O. INITIALLY, THEN 300 MG (1 TAB) IN 6 HRS, AND QD FOR 2 DAYS

**PLUS**

- PRIMAQUINE BASE 30 MG (2 TABS) P.O. PER DAY FOR 14 DAYS

# TREATMENT

- UNCOMPLICATED  
CHLOROQUINE RESISTANT  
INFECTIONS:
  - QUININE 650 MG PO TID X 3 DAYS  
AND DOXYCYCLINE 100 MG PO BID  
X 7 DAYS

## COMPLICATED OR SEVERE INFECTIONS:

- Intravenous antimalarial medications

# TREATMENT

- VIVAX AND OVALE THERAPY SHOULD INCLUDE PRIMAQUINE 30 MG BASE PO QD X 14 DAYS

# OPTIMAL TREATMENT APPROACH

- RAPID CASE IDENTIFICATION
- RAPID PARASITOLOGICAL CLASSIFICATION
- RAPID INITIATION OF THERAPY
- RAPID INITIATION OF SUPPORTIVE CARE

# CONTROL OF MALARIA

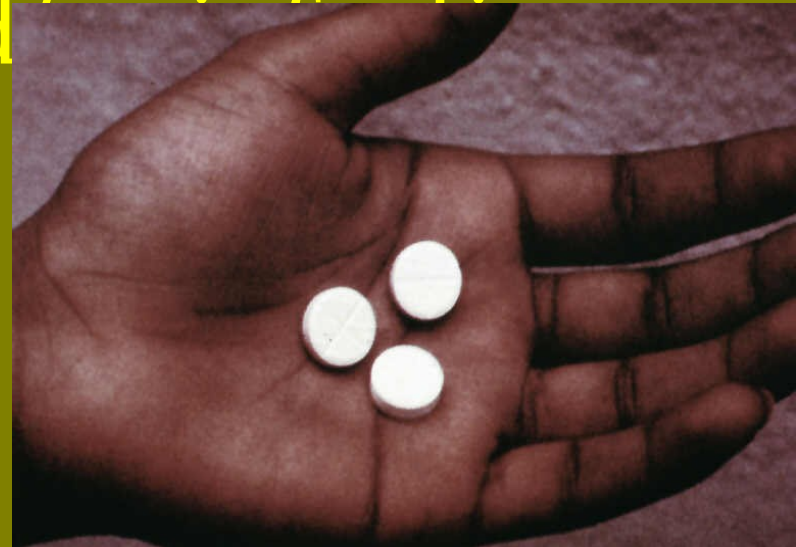
- GLOBAL ERADICATION EFFORTS BY WHO IN 1950s
- EFFORTS NOW FOCUS ON CONTROL vs ERADICATION

# POINTS OF ATTACK

1. ATTACK THE PARASITE IN THE HUMAN HOST
2. REDUCE CONTACT BETWEEN HUMANS AND MOSQUITOES
3. DECREASE MOSQUITO POPULATION

# ATTACK THE PARASITE IN THE HUMAN HOST

- Treat malaria infections with effective medications
- Use prophylactic drugs to prevent illness and transmission





# ATTACK THE PARASITE IN THE HUMAN HOST

- CHEMOPROPHYLAXIS BASED ON CURRENT DRUG RESISTANCE PATTERNS
- MEFLOQUINE FIRST LINE PROPHYLAXIS
  - Mefloquine 250mg. po q week, 2 wks prior to exposure and for 4 wks after exposure
- DOXYCYCLINE AS SECOND LINE DRUG
  - Doxy 100mg. po qd, 2days prior to exposure and for 4 wks after exposure
- Primaquine 30 mg. po qd x 14 days terminal prophylaxis



# REDUCE CONTACT BETWEEN HUMANS AND MOSQUITOES

- PERSONAL PROTECTIVE MEASURES
  - PROPER WEARING OF UNIFORM
  - DEET LOTION
  - PERMETHRIN TREATED UNIFORMS
  - BED NETS





# DECREASE MOSQUITO POPULATION

- ♦ **SURVEILLANCE OF MOSQUITO POPULATIONS**
- ♦ **ID AND ELIMINATION OF BREEDING SITES**
- ♦ **PROPER INSECTICIDE APPLICATION**
  - **ATTACK LARVAL STAGES**
  - **ATTACK ADULT MOSQUITO**



# SOMALIA



◆ 48 CASES IN COUNTRY

◆ 83 CASES FOLLOWING  
RETURN

-62 USA, 21 USMC





# SOMALIA

- ISSUES

- COMMAND RESPONSIBILITY
- COMPLIANCE: SWITCH TO MEFLOQUINE
- LOCATION OF CAMPS
- WEATHER → POOR USAGE OF DEET
- NO USE OF BED NETS
- PRIMAQUINE TERMINAL PROPHYLAXIS
  - USA HAD NOT RECOMMENDED

# SOMALIA CONCLUSIONS



♦CASES RESULTED FROM FAILURE TO IMPLEMENT PROPER PROPHYLAXIS AND PERSONAL PROTECTION

♦MEFLOQUINE MORE EFFECTIVE THAN DOXYCYCLINE IN USMC



# COMMAND RESPONSIBILITY

- Field Marshall Sir William Slim:  
“If the overall result was less than 95% positive, I sacked the commanding officer. I only had to sack three; by then the rest had got my meaning.”
- Sir Neil Cantlie, Dir Gen Brit Army Med Services  
“When for the first time in history a combatant officer was considered unfit to command a unit on the grounds that he had allowed his men to become ineffective through disease, a new day in military medicine dawned. The clouds of forgetfulness must not be allowed to overshadow the brightness of that day.”





# CONCLUSIONS

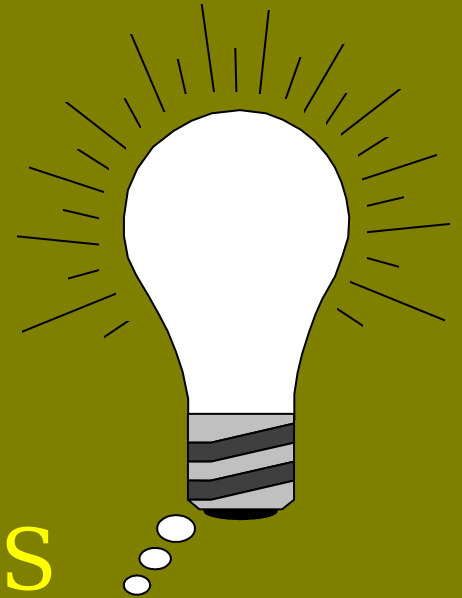
EDUCATION AND AGGRESSIVE  
MONITORING OF COMPLIANCE  
NEEDED FOR CHEMOPROPHYLAXIS  
AND PERSONAL PROTECTIVE  
MEASURES TO WORK



# SUMMARY

- MALARIA DRUG RESISTANCE IS INCREASING
- GREAT IMPORTANCE OF **PERSONAL PROTECTIVE MEASURES**
- AGGRESSIVE MONITORING NEEDED TO ENFORCE PPM AT COMMAND LEVEL
- DX REQUIRES HIGH INDEX OF SUSPICION
- REGARD AND MANAGE MALARIA AS MEDICAL EMERGENCY

# MALARIA!



- REMEMBER:

FLU-LIKE SYMPTOMS

+

‘RECENT’ HX TRAVEL  
TO MALARIOUS AREA

=

THINK MALARIA